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Inferring cellular forces from image stacks

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Non-technical Summary

The importance of cellular forces to a wide range of developmental and disease processes is widely recognized, but measuring these forces is challenging. Measurements are especially difficult in 3D, where cells take polyhedron-like shapes, can be buried beneath other cells, and can have face-specific tensions that change over time and that drive cell movements and form precise multi-cellular structures. Here we present a computer-based technique that infers these forces from cell images, and in 8-cell mouse embryos does so with errors smaller than 10%. The technique has the potential to shed light on the mechanics of cellular processes ranging from embryogenesis to cancer metastasis.

Abstract

Although the importance of cellular forces to a wide range of embryogenesis and disease processes is widely recognized, measuring these forces is challenging, especially in 3D. Here, we introduce CellFIT-3D, a force inference technique that allows tension maps for 3D cellular systems to be estimated from image stacks. Like its predecessors, Video Force Microscopy and CellFIT, this cell mechanics technique assumes boundary-specific tensions to be the primary drivers, and it constructs force-balance equations based on triple-junction dihedral angles. The technique involves image processing, segmenting of cells, grouping of cell outlines, calculation of dihedral planes, averaging along 3D triple junctions, and matrix equation assembly and solution. The equations tend to be strongly overdetermined, allowing indistinct triple junctions to be ignored and solution error estimates to be determined. Application to clean and noisy synthetic data generated using Surface Evolver gave tension errors of 1.6 to 7%, and analyses of 8-cell murine embryos gave estimated errors smaller than 10%, the uncertainty of companion aspiration experiments. Other possible areas of application include morphogenesis, cancer metastasis and tissue engineering.

Keywords

Cell mechanics, force inference, embryogenesis, murine embryos, CellFIT-3D, CellFIT, Video Force Microscopy (VFM), Surface Evolver.

Cellular forces

The importance of cellular forces to morphogenesis, wound healing and disease is now widely recognized [1-5]. Experimental studies have shown that irregularities in these forces can give rise to developmental defects and other malformations [6], and computational models have shown that alterations as small as 20% can affect clinical outcomes [7].

In many settings, these driving forces can be treated as equivalent interfacial tensions γ_i along cell-cell and cell-medium boundaries [8,9] (**Supplementary Fig. 1**), a concept considered as early as the 1960s [10,11] but quickly dropped in favor of the differential adhesion hypothesis [12]. Computer simulations of sorting and other cell movements carried out by the authors, however, showed that a wide range of cell and tissue movements were in fact driven largely by interfacial tensions [13-15] (to which cell-cell adhesions make a counteracting contribution [8]), leading to a new paradigm [2,16-20]. Information about these tensions can be obtained through a variety of experimental techniques, that in generally decreasing scale include: pipette aspiration, deformable substrates, engineered droplets, laser ablation, atomic force microscopy, optical tweezers, magnetic cytometry, and FRET [18,21-31]. Unfortunately, these techniques can be expensive, time consuming, invasive, or destructive, and all except substrate deformation, FRET and droplet methods provide information about only a single cell surface at one moment in time.

When force inference techniques for 2D systems [32-36] entered the scene in 2010, they made spacio-temporal maps of cellular forces possible for any available image, including historical ones. Force inference techniques, like CellFIT which is here denoted as CellFIT-2D to distinguish it from the present 3D version, are based on two primary mechanical assumptions: that the tension-carrying boundaries that meet at any triple junction (TJ) [36] produce mechanical equilibrium there, and that these tensions are constant over the span of any given boundary or interface (**Supplementary Movie 1**). Force inference techniques in this class require only that cell edges be visible, they involve no mechanical intervention, and they produce no additional damage beyond that caused by microscopy. Data from CellFIT-2D revealed tension variations around the perimeters of individual cells, differences between cells in a single population and between populations, elevated tensions along inter-population boundaries [36], and purposeful temporal variations [17].

On the strength of this and other evidence, we propose that cells move and organize primarily by gradual and carefully-orchestrated changes in interfacial tensions and protrusion contractions, with the latter acting along the interface between cells and serving as a special case of these tensions. Finite element studies have shown that force imbalances at triple junctions cause those junctions to move until TJ angles produce an equilibrium configuration or until a high-tension boundary shortens enough to produce a neighbor change [15]. If these forces were to change systematically and gradually with time, a series of carefully-controlled cellular and tissue morphologies could be produced. Modeling studies of whole embryos [37] found that physiological motions were best matched by time-varying driving forces, but the reason for this finding was unclear when those studies were carried out. The finding was contrary to the terraced nature of these forces tacitly assumed by modelers and biologists. In the context of embryogenesis, the context in which the most active research on cell mechanics and morphogenesis was occurring, the underlying mindset was that gene expression or some other process set up conditions for the next developmental step and then it moved ahead on the basis of those conditions. The revised understanding for which we here argue, is that regulatory networks and

mechanobiological circuits play a much more active role in controlling the driving forces from moment to moment. If cellular forces do indeed change with time as these studies suggest, then techniques like CellFIT – which can take snapshots of the forces in all of the cells visible in a field of view and do so at multiple sequential times – become particularly useful.

Other kinds of forces – such as intracellular pressures and deviatoric stresses associated with cytoplasm deformation – could be included, as was done in Video Force Microscopy (VFM) [17]. Line tensions along the curvilinear interfaces between trios of cells could also, in principle, be included, but we suspect that their contribution is minimal. The picture of gradual force changes we present should not be confused with models in which increasing forces are applied to elastic systems. Elastic systems would spring back to their initial state should the driving forces be eliminated, though real embryos and their tissues do not. Instead, we contend that embryos and their tissues behave primarily in a plastic manner [38], with their morphogenetic deformations being essentially irreversible due to cell neighbor changes. An apparent exception to this statement is the small elastic component of deformation that can occasionally be observed, as when momentary forces are applied, and a reversible response occurs only because cells do not have time to change neighbours and lock the temporary geometry in place. We have argued elsewhere [28] that viscous, visco-elastic and elastic tissue behavior can all be produced by cells that have constant edge tensions and viscous cytoplasm.

Interestingly, the triple-junction force balances on which CellFIT is based do not depend on the mechanical characteristics of the cell membrane system, including whether it is elastic, viscous, governed by rate constants, affected by endocytosis, influenced by adhesion systems or altered by cortical components. In a sense, CellFIT operates one level up from these important details, and simply provides the total relative tension acting along any particular interface, without regard to how it is generated. In addition, intracellular pressures and any stresses from viscous or contained elastic components act primarily normal to the interface (except perhaps during laser ablation experiments, which violate other CellFIT assumptions) and likewise, do not affect the triple-junction force balances. These fortuitous circumstances make CellFIT applicable to a very wide range of biological applications. CellFIT would not be applicable, however, to interfaces where cells are adhered to a substrate, because the forces carried by that substrate would generally not be known. If such forces were known, however, governing equations that include them could be constructed [5], and a suitably-adapted version of CellFIT applied. It would also not be applicable to interfaces that contain significant spatial force variations along individual cell faces. One might hope that the quality measures that form part of CellFIT would aid in identifying situations of this kind.

In this article, we outline a series of steps (**Fig. 1**) that allow CellFIT-2D to be extended so that cellular forces in 3D aggregates and tissues can be inferred from serial sections. When applied to synthetic sections generated using Surface Evolver (SE) [39], the tensions calculated by CellFIT-3D had errors as small as 1.6%. CellFIT-3D was then applied to 8-cell compaction-stage mouse embryos, and the tensions found had estimated errors lower than the 10% uncertainty associated with accompanying aspiration experiments.

CellFIT-3D

As in the finite element models we have used to study embryonic cells for many years and in VFM and CellFIT-2D, we here assume that the sub-cellular forces generated by various structural protein and

adhesion systems can be deemed to generate an equivalent tension tangent to the cell membrane [13,15,17,36,40] (**Supplementary Fig. 1**). The total tension along any given cell-cell or cell-membrane interface i is denoted γ_i , and it is assumed to be specific to that interface, whether an edge in 2D or a face in 3D [41].

Furthermore, whenever three cells meet at a particular point and a cutting plane is constructed normal to the TJ between those cells at that point, the vector sum of the membrane tensions in that plane must add to zero. Studies of 2D cellular systems showed that the membrane angles at TJs are unaffected by intracellular pressures or deviatoric stresses associated with reshaping of the viscous cellular cytoplasm. Consequently, TJs can be analyzed and used to calculate membrane tensions γ_i without reference to pressure or viscous forces.

Cellular pressures were not calculated as part of CellFIT-3D, but one could presumably calculate them, as is done in 2D, after the interfacial tensions γ_i are determined. In 3D, the local principal curvatures k_1 and k_2 of cell-cell or cell-medium interfaces change with position unless they are spherical. However, the mean curvature $(k_1+k_2)/2$ would be constant throughout any one of these surfaces if it carries a constant isotropic tension γ_i , and the pressure difference ΔP_i across it would be given by the Young-Laplace equation

$$\Delta P_i = \gamma_i (k_1 + k_2). \tag{1}$$

To determine the spatially-varying principal curvatures or even the mean curvature of a surface based on cuts through it, however, is a problem beyond the scope of the present study. For the present, there is probably no reason to calculate intracellular pressures or their differences, as their relevance to development is still unclear and methods to verify them experimentally are limited.

Because of the geometric complexity of 3D systems, it is valuable to distinguish between the curvilinear triple junctions or triple edges (TEs) that arise between trios of contacting cells or between pairs of cells and the medium, and the points at which those junctional curves pass through individual confocal or other sections. We use the term “triplet” to describe the point where a TE passes through a section and three, or occasionally more, cell boundaries are seen to converge (as in the boxed area of **Fig. 1a**). The cell membranes immediately adjacent to the point are considered part of the triplet.

A series of image processing and geometric and mechanical analyses steps are required to implement CellFIT-3D, and they are outlined here.

Image capture. As we will show, the quality of a CellFIT-3D analysis depends on image resolution and section spacing. Reliable tension information can be obtained from data like **Fig. 1a**, a portion of a 512 by 512 pixel image from a confocal stack of an 8-cell murine embryo. There were approximately 125 pixels across the diameter of a typical cell, and individual cells were transected an average of 18 times as a result of the 2 μ m section spacing.

Image enhancement. The cell boundaries in these images may seem well defined, but at the pixel level (**Fig. 1b**) where computational algorithms work, noise, gaps and other anomalies become apparent. An ideal image would have accurate, gap-free white outlines a single pixel wide and a pure black background. A number of image processing algorithms for amending the images toward this ideal were appraised, and an edge enhancing coherence filter [42] that smoothed the boundaries (**Fig. 1c**) and

largely closed the gaps (**Fig. 1d**) was selected. Relevant computer code for this step and a number of the others, as well for as the validation procedures, is available through the indicated references.

Image segmentation. Cells in the processed images were segmented (**Fig. 1e**) using a watershed algorithm called SeedWater [43,44]. The algorithm treats the bright pixels at cell edges as if they defined mountain range heights and it begins to flood each mountain-surrounded region from a relatively dark pixel or low-lying point called a seed point. The software chooses these seed points, but users can manually adjust them. Boundaries are found indirectly, based on where the flooded zones contact each other or where they end at the outer mountain ranges. This approach is less sensitive to intracellular noise – spurious hills in low lying regions – and partial gaps than are standard edge tracing techniques. The resulting boundaries may, however, still contain pixel-scale positional noise (**Fig. 1f**) and other imperfections.

Grouping of triple edges. In 3D, individual triple edges (TE) may appear as triplets in multiple images and it is necessary to associate these occurrences with each other. As with boundary identification, it is preferable to work with cell cross-sectional areas, and we use a cell-grouping algorithm that considers area overlap across adjacent images, centroid collinearity and other geometric features [45] to group cross-sections associated with the same cell (**Fig. 1g**) and in turn to identify triplets belonging to the same TE. If sections are spaced sufficiently close, seed-point proximity can be used to group the outlines. Grouping also makes possible 3D cell models for visualization and verification of mesoscale geometry and topology.

Mesh generation. The pixelated boundaries of sectioned cells may have complex shapes with reversing curvature and may contain noise. In order to obtain good triplet approach angles, and informed by CellFIT-2D [36], we fit cubic splines to the boundary pixels along each edge and use these splines to generate uniformly-spaced mesh points (**Fig. 1h**).

In-plane angles. CellFIT-2D showed that accurate angles are crucial to the tension calculations, and we found that membrane end-point directions could be estimated well by fitting separate circular arcs to the last 4 or 5 mesh points on each end (**Fig. 1i**), as in 2D [36]. Standard software exists for arc fitting, the fit is coordinate indifferent, it provides a second-order approximation to the shape of the boundary terminus, and it attenuates noise.

Triplets that appeared in at least 3 successive images were identified algorithmically, and shown graphically in their corresponding sections (**Fig. 1j**). This strategy was more efficient than manually vectorizing triplets of interest [46]. The graphic consisted of a circle at the calculated triplet location and three vectors with circles at their ends in the calculated approach directions. The graphics were automatically overlaid on the raw images and manual angular adjustments were made as needed (**Fig 1k**).

Local dihedral angles. To convert the in-plane angles defined by these graphic triplets to true dihedral angles, splines (shown as orange curves in **Fig. 1l**) were constructed through sets of grouped triplets. Construction of a reliable spline required a TE to appear in at least three images. Local dihedral planes (**Fig. 1m**) were constructed normal to the spline at each triplet, and the boundary vectors were mapped onto them (white arrows) by projection normal to the planes. Finally, a pair of equilibrium equations was constructed for two arbitrary orthogonal directions in each plane. These equations define the ratios that the three boundary tensions γ_i must have for that section of the TE to be in equilibrium.

Average dihedral angles. The equation pairs generated at various triplets along any one TE can give different force ratios, and the quality of the equations varied. For example, if an image plane is strongly oblique to a TE, the fluorescent signals from one or more of its boundaries may be spread out and grainy. In addition, errors in calculated boundary directions can amplify when projected onto highly-tilted dihedral planes, making information from these planes less trustworthy. When more than 3 equation pairs are available, they are automatically checked for consistency and equations from the ends of the spine, where the dihedral planes tend to be more tilted, eliminated if appropriate. Increasing the number of sections per cell improves the geometry of the spline and increases the number of available equation pairs. However, even when there are 7 slices per cell on average (orange curve in **Supplementary Fig. 3**), approximately two thirds of the TEs still appear in two or fewer sections. To amalgamate the multiple equation pairs associated with any one TE into a single set of ratios $\gamma_i:\gamma_j:\gamma_k$, we recommend a least-squares ratio solver [47].

In contrast to these steps, one could construct point clouds from the meshing points associated with each boundary. One could then fit mathematical surfaces to each cloud, or a portion thereof, and use mathematical descriptions to calculate the TE contact angles. Unfortunately, the shapes that arise are complex, often having reversing concavity and other challenging features, making this approach impractical [48]. One could also take in-plane angles as surrogates for dihedrals and apply CellFIT-2D to individual sections. However, doing so ignores the oblique angles of typical TEs to these sections, and produces tension errors of 50% [48]. Alternatively, for specific TEs, one might use the in-plane angles of the triplet apparently most normal to that TE [21], but numerical tests show that even this approach introduces approximately 5% additional error in the dihedral angle equations, in addition to foregoing the statistical benefits of multi-triplet averages.

Assembly and solution of tension equations. The previous step produces either a pair of equilibrium equations or two tension ratios for each of the n TEs having a sufficient number of useable triplets. In either case, these equations can be assembled into a single matrix equation of the form

$$\mathbf{G} \boldsymbol{\gamma} = \mathbf{0}, \tag{2}$$

where \mathbf{G} has $2n$ rows and m columns, and m is the number of tensions γ_i that appear in one or more of the $2n$ force-balance equations, and whose values will in due course be found. These equations correspond exactly to the equation pairs that arise in CellFIT-2D [36], and their assembly, solution and evaluation are identical, unaffected by the dimensionality of the host space.

A unique solution to this homogeneous and overdetermined system can be found by constructing and solving the least-squares system

$$\begin{bmatrix} \mathbf{G}_\gamma^T \mathbf{G}_\gamma & \mathbf{C}^T \\ \mathbf{C} & \mathbf{0} \end{bmatrix} \begin{Bmatrix} \gamma_1 \\ \vdots \\ \gamma_{N_{Tensions}} \\ \lambda_1 \end{Bmatrix} = \begin{Bmatrix} 0 \\ \vdots \\ 0 \\ N_{Tensions} \end{Bmatrix}, \tag{3}$$

or

$$\mathbf{G}^* \boldsymbol{\gamma}^* = \mathbf{n} \tag{4}$$

where

$$\mathbf{C} = \{1 \dots 1\}, \quad (5)$$

a choice that selects the particular set of scaled γ s having a unit mean.

Solution evaluation. The toolkit developed for assessing CellFIT-2D solutions was found to apply equally well to CellFIT-3D. The overall quality of the equations is assessed using the condition number of \mathbf{G}^* , a value equal to the ratio of its largest and smallest eigenvalues. This ratio portends the degree to which error in \mathbf{n} is amplified in γ^* [49], and so indicates the sensitivity of the calculated tensions to angular and other types of error. The condition number will vary with matrix size, and magnitudes substantially higher than those shown here and in other CellFIT analyses [36] may signal structural problems with the equations and reduced solution accuracy.

In contrast, the solution residual

$$\mathbf{r} = \mathbf{G} \gamma, \quad (6)$$

provides a measure of how well the pair of equations associated with each TE is satisfied by the least-squares solution γ . The paired components of \mathbf{r} indicate the degree to which each TE is out of balance and they are useful for identifying triplets that may have been inaccurately placed during the automated meshing steps and may require manual adjustment. Finally, the scaled cofactors of \mathbf{G}^* , also known as standard errors [36], indicate the confidence levels associated with individual tensions.

Solution reporting. The calculated tensions can be reported in many different forms, including that shown in **Fig. 1n**. As with CellFIT-2D, display of residuals or confidence levels [36,50] might be useful, but doing so in 3D is more challenging.

Validation

To assess the CellFIT-3D algorithms, they were tested on synthetic sections generated using Surface Evolver [51] (**Fig. 2** and **Supplementary Material**) for which ground truth tensions were known. The synthetic data were designed to approximately match the 18 slices per cell of the murine embryo data used here, and the images data were processed as outlined in **Fig. 1**. The error in the calculated tensions was found to depend on slice spacing and image resolution, but was insensitive to the number of cells analyzed, whether those cells were surrounded by medium or cut cells, and the range of the tension values γ_i within the model.

Increasing the number of slices per cell increases the number of dihedral planes per TE (**Supplementary Fig. 3**) and allows strongly-sloped planes near their ends to be ignored. Seven slices per cell was found to give tension errors of 3.2% in isolated aggregates of 8 cells, while 14 slices per cell, often not difficult to obtain experimentally, reduced that error to only 1.6% (**Fig. 2c**). In all cases, the number of equations was more than adequate to solve for the unknown tensions (see **Supplementary Material**).

Portions of isolated synthetic aggregates of 8 and 50 cells were eroded away in order to generate a range of aggregate sizes and conditions – from those fully surrounded by medium, to those with mixed medium and cut cell edges, to fragments containing no complete cells. For aggregates with 14 slices per

cell, the errors depended primarily on the image resolution, with 400 and 200 pixels per cell diameter generating errors of 3% and 8%, respectively.

Error was then added to the averaged dihedral angles, and **Fig. 3** shows how adding angular noise to the average dihedral angles affects tension error, residual and standard error. The angular noise was randomly assigned using a Gaussian distribution. Conveniently, all three track similarly with noise, allowing the latter two – which can be calculated from the governing equations associated with any given data set – to indicate the likely angular error (input noise) and tension error (output uncertainty). A mean angular noise of 5 degrees, a value consistent with typical manual digitizing errors of 5° [36], for example, caused tension errors of 7.1% and residual and standard errors of 9.4% and 5.2%, respectively.

Application to murine embryos

CellFIT was then used to investigate 8-cell murine embryos (**Fig. 4a** and **Supplementary Materials**) undergoing compaction, a process considered to be driven by surface tensions [21,52]. Interfaces were typically seen in at least 5 slices. Some manual adjustments were necessary in places where the automated meshing was inaccurate. For Time A, 90 minutes after the last 4-cell stage blastomeres divided, the CellFIT-3D tensions had residuals of 0.031 and standard errors of 5.4% (**Fig. 4b**), while the values for Time B (**Fig. 4c**), 150 minutes later, were slightly higher. The corresponding boxes in **Fig. 3** suggest input errors corresponding to noise levels between 2 and 6 and tension errors of 3-9%.

The surface tensions of all 8 cells (**Fig. 4a**) were measured by pipette aspiration [21] and the experimental error estimated from repeated tests was 10%. Measurement of tensions on interior surfaces is more challenging, but could be done using optical tweezers or laser ablation, though the latter is destructive and precludes the taking force measurements at multiple times or locations. The RMS errors between the CellFIT-3D and aspiration values at Times A and B (**Figs 4b and c**) were 14.6% and 11.6%, respectively, values consistent with aspiration errors of 10% and standard errors of 5.4% and 6.4%, for Times A and B, respectively. The surface tensions of individual cells changed between Times A and B, and the time delay between sequential aspiration measurements may have contributed additional error. In contrast, CellFIT typically draws its data from the much narrower window of time required to collect a single set of sections. Several other 8-cell murine embryos were analyzed with CellFIT, and similar results were obtained.

Discussion

This article demonstrates that cellular forces (interfacial tensions) in 3D aggregates can be inferred from serial sections. In principle, any image sources including confocal sections or histological sections could be used, provided that cell boundaries are well defined and there are at least 6 to 8 slices per cell. Furthermore, tension errors can in principle be as low as 1.6%, considerably lower than the variability of typical experimental tensometry techniques. Because the assembled CellFIT equations are overdetermined, input and solution quality can be assessed using residuals and standard errors, and TEs with apparently discordant angles or other anomalies can be identified and removed without jeopardizing the tension inference process. These quality measures could also serve to identify situations where membrane tensions are not uniform as assumed in CellFIT, as could be the case should

undulating membrane geometries arise, or localized tension sources give rise to nonuniformities in membrane tension.

CellFIT-3D builds on its 2D counterpart, and its three primary challenges – obtaining cell outlines, connecting triplets in successive images, and calculating true dihedral angles – can be overcome using relatively straightforward algorithms. Like its counterpart, CellFIT-3D provides only tension ratios, and should scaled values be required, direct force measurements must be obtained. Fortunately, these ratios are often all that is required [21], as it is these ratios that govern cell shapes and motion patterns, with cytoplasm viscosity and force strength determining motion rates, only [53].

Force inference methods are possible because the relative angles at TEs in equilibrium depend uniquely on membrane forces, even though other mesoscale features (see **Supplementary Text**) do not [54,55]. Should four or more cell edges impinge on a single junction or there be a rosette [56], the number of unknowns exceeds the number of equations by more than one, and a unique tension ratio does not exist. However, if the equations and unknowns associated with such junctions are assembled with those from other suitably-connected TEs, a unique and trustworthy solution should still be possible [36]. In principle, CellFIT-3D could be extended to calculate intracellular pressures, as in CellFIT-2D [36], but calculating the required surface curvatures is much more difficult in 3D, there is currently little experimental interest in these pressures, and their consequences for cell movement remain unclear. The effects of viscous cytoplasmic forces could also be incorporated easily, with their contributions to TJ force balances calculated in a manner similar to that used in VFM.

If the cellular forces that drive cell movements, mesoscale assembly and bulk tissue motions turn out to be time-varying, as recent evidence suggests, then techniques like CellFIT that can provide force maps over a whole field of view and at closely spaced times become particularly useful. Hopefully, in time, the technique presented here will provide new insights into the movements of single or small groups of cells during morphogenesis and diseases like metastasis, formation of mesoscale structures such as acini in various organs and engineered tissues, and bulk tissue movements.

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Authors' Contributions

The force inference concepts were developed and implemented by GWB, JHV and AE; the experiments were conducted by JLM in the lab of TH; the Surface Evolver models were developed by SC and JHV; the paper was written primarily by GWB.

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Competing Interests

We have no competing interests.

For Review Only

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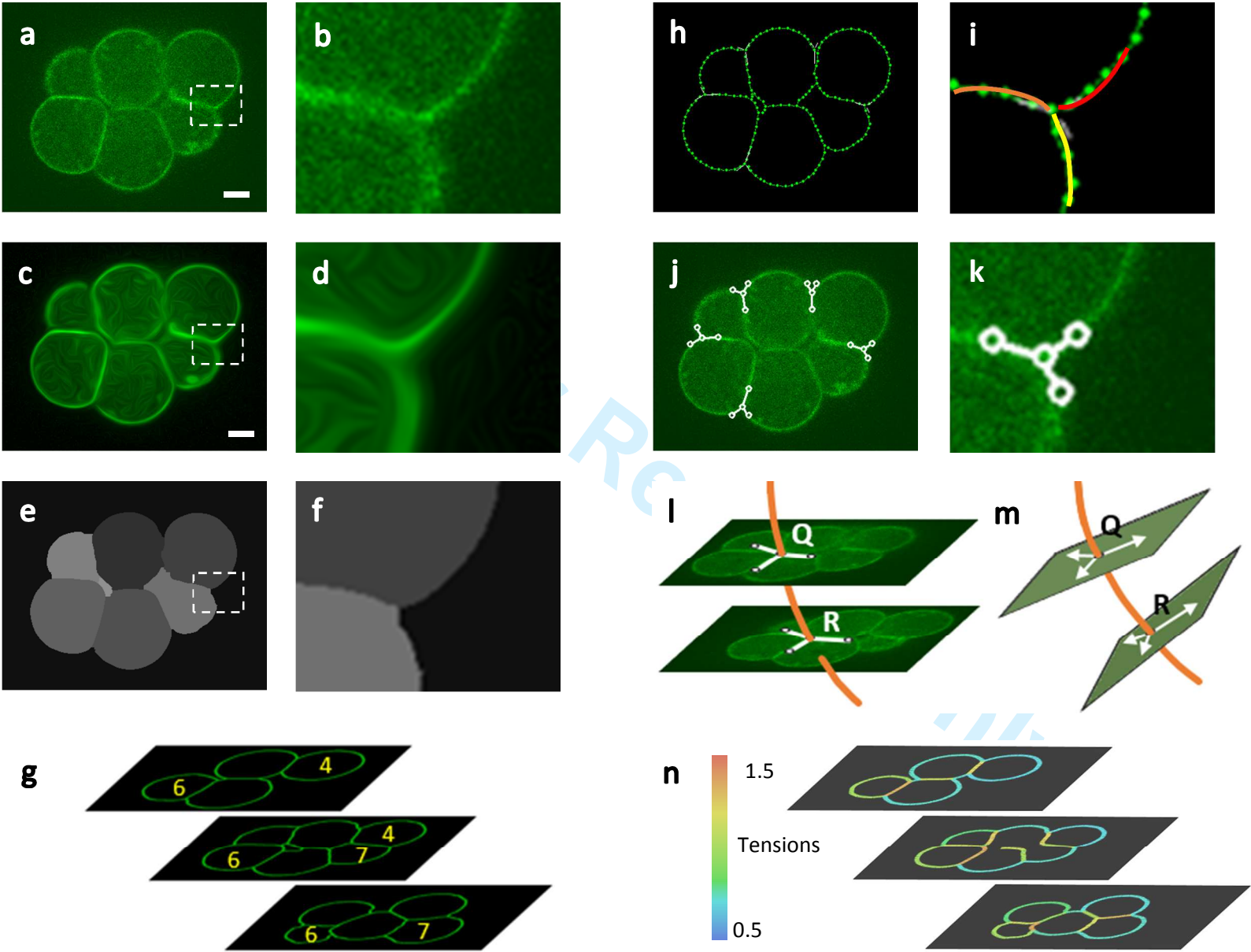
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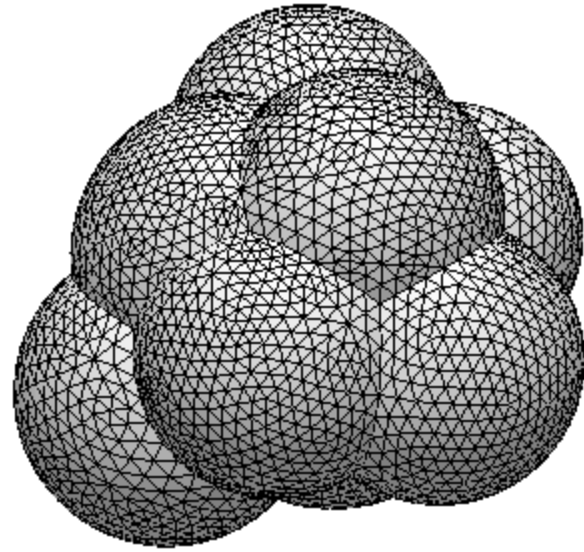
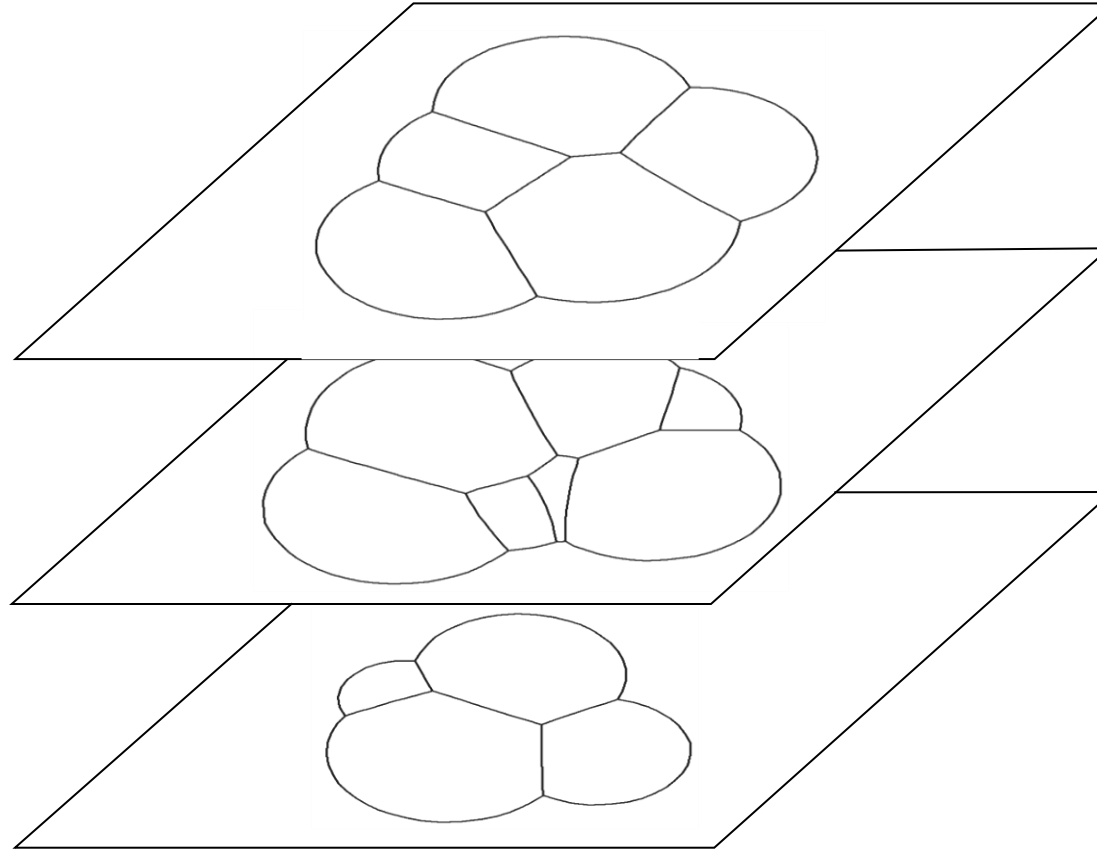
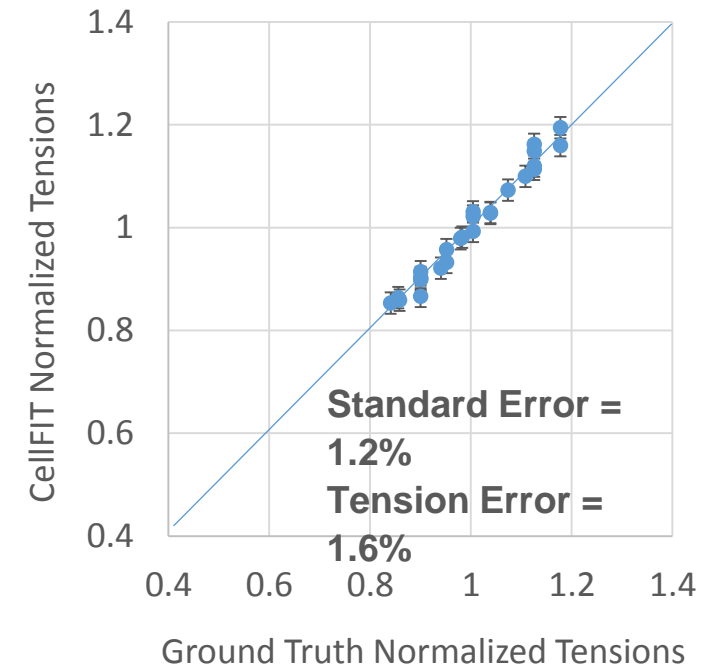
Figure 1 - An overview of CellFIT-3D. A raw confocal image of an 8-cell stage murine embryo is shown in **a** (scale bar = 10 μm), while **b** shows an enlargement of the boxed area. An edge enhancing coherence filter [42] was used to smooth the boundaries **c** and close their gaps **d**. Parameters used: Scheme, implicit discretization; total diffusion time = 25; Gaussian sigma = 3. SeedWater [43]-segmented cells are visible in **e** and **f**, each denoted by in a different intensity of grey. Corresponding cells in successive images were grouped together **g** and TEs that appeared in multiple images identified. A fine mesh **h** was constructed along each cell boundary and its approach angle to any given triplet determined by circular arc fitting **i**. Tangent vectors **j** were calculated automatically and adjusted manually as needed **k**. Splines were fit through the triplets that appeared in successive images **l**, and dihedral planes and angles calculated **m**. Least squares equations were constructed and solved (see text) and calculated tensions displayed **n**.

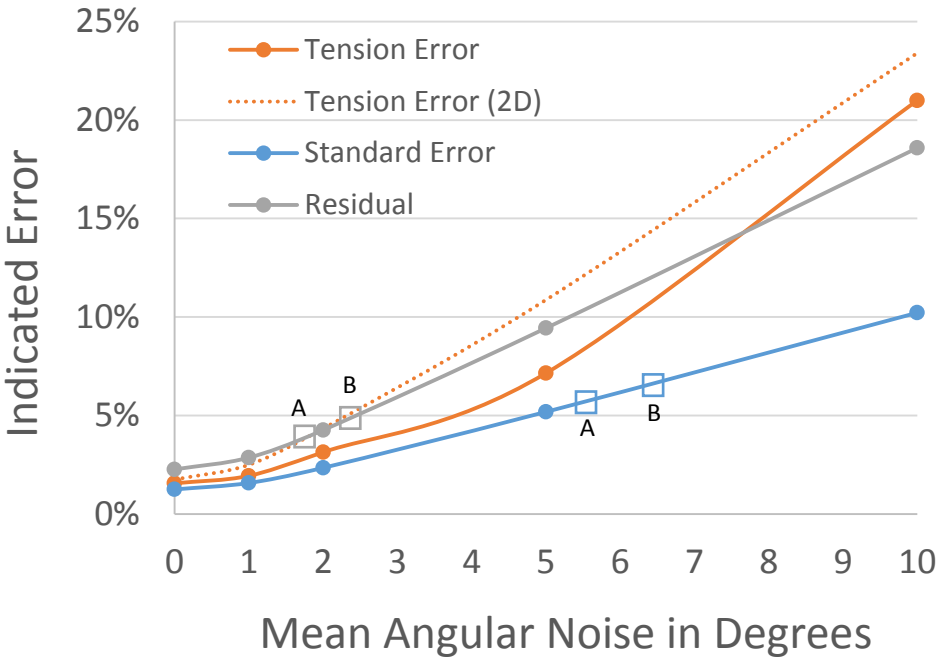
Figure 2 - Synthetic data and analysis. A model with 8 cells and randomly assigned interfacial tension values varying from each other by 30% was generated using Surface Evolver **a**, and used to create synthetic sections **b**. The tensions calculated by CellFIT-3D are shown in **c** and error bars indicate their associated standard errors. The RMS tension error relative to ground truth was 1.6%.

Figure 3 - Noise response. Tension error grows as mean angular noise, of the amounts indicated in angular degrees, is introduced into the averaged dihedral angles. The standard error and residual behave similarly, as does the tension error in a 20-cell epithelium analyzed using CellFIT-2D [36]. The standard errors and residuals associated with murine embryo analyses (**Figs 4b** and **c**) are shown as boxes along their corresponding curves, and they indicate, respectively, equivalent angular noise in the averaged dihedral angles of 6-7% and 2-3%. Tension errors can in turn be deduced from these noise levels to be no greater than 7% (see **Supplementary Text**).

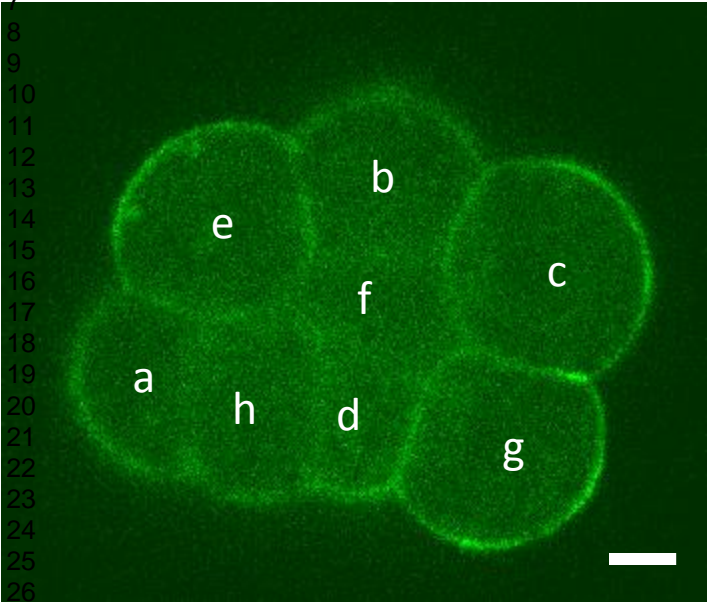
Figure 4 - Analysis of murine embryos. **a**, Compaction-stage murine embryos consisting of 8 cells were imaged such that cells were nominally 120 pixels in diameter, and there were on average 18 sections through each cell (scale bar = 10 μm). Tensions measured using micropipette aspiration at Time A (90 minutes after the last division from 4- to 8-cell stage) are shown in **b** and those at Time B, 150 minutes later, in **c**. CellFit-3D inferred tensions have been scaled so that their average value is the same as that of the values from the pipette aspiration experiments. Error bars on the experimental values are $\pm 10\%$ (see text) while CellFIT-3D tensions show their respective Standard Errors. In this figure, only, Tension Errors are with respect to aspiration experiments not ground truth.



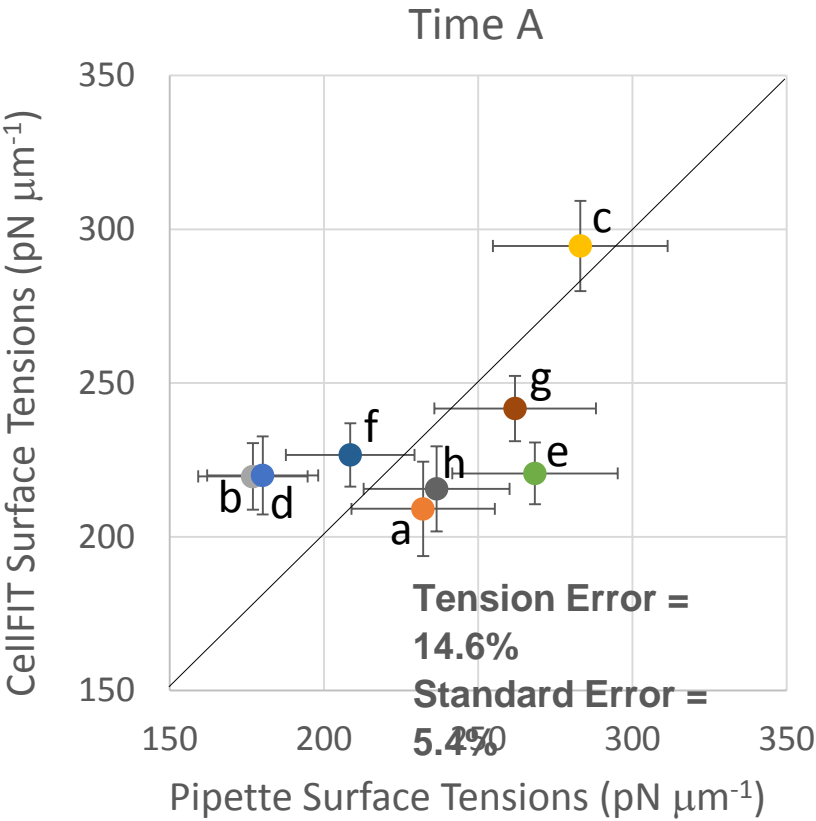
**a.****b.****c.**



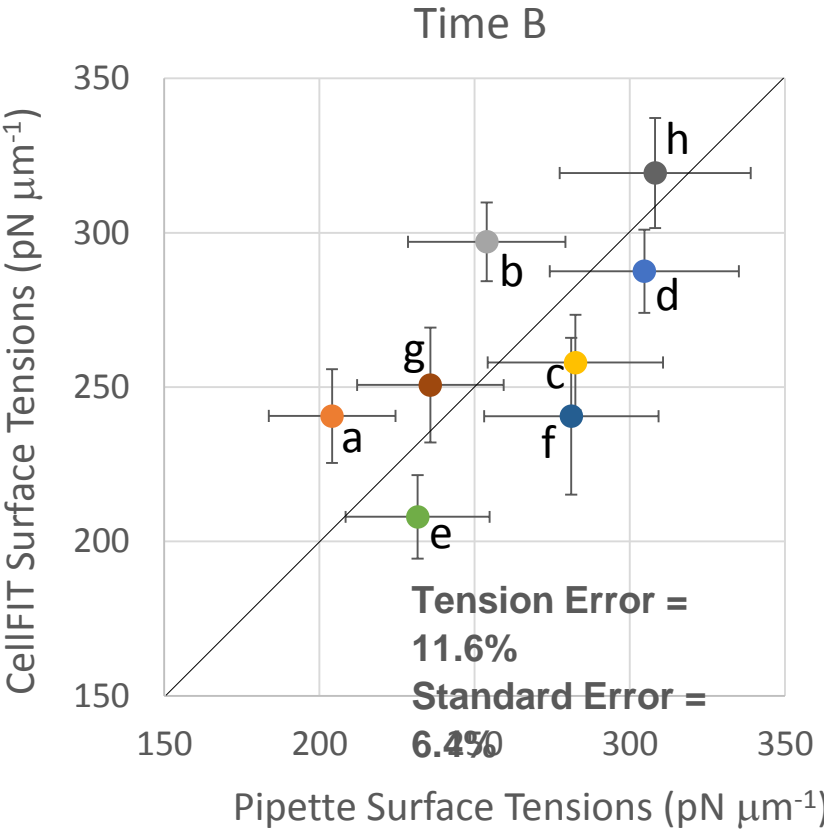
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a.



b.



c.